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Sampling Strategies Near a Low-Order Detonation and a Target at an Artillery Impact Area

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Charles A. Ramsey, Dennis J. Lambert, Kevin L. Bjella,
and Nancy M. Perron

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 (SERDP)

ABSTRACT

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PREFACE

This report was prepared by Dr. Thomas F. Jenkins, Alan D. Hewitt, and Kevin L. Bjella, Environmental Sciences Branch, U.S. Army Engineer Research and Development Center (ERDC), Cold Regions Research and Engineering Laboratory (CRREL), Hanover, New Hampshire; Thomas A. Ranney, Science and Technology Corporation, Hanover, New Hampshire; Charles A. Ramsey, EnviroStat, Inc., Fort Collins, Colorado; Dennis J. Lambert, Engineering Resources Branch, ERDC-CRREL; and Nancy M. Perron, Snow and Ice Branch, ERDC-CRREL.

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1 INTRODUCTION

A series of papers has been published describing the difficulty in collecting soil samples representative of the mean analyte concentrations at areas contaminated with residues of energetic compounds (Ampleman et al. 2003a, b; Jenkins et al. 1996, 1997, 1999, 2001, 2004; Pennington et al. 2001, 2002, 2003; Thiboutot et al. 1998, 2003; Walsh et al. 2001, 2004). This difficulty is because residues often exist as particulates and are distributed heterogeneously on the surface. Because such particulate residues may serve as the major source of potential off-site migration of these compounds, it can be important to estimate the mass of energetic materials present in these surface soils.

There are a number of different types of DoD training ranges where various types of munitions are used. These include artillery and mortar range impact areas, bombing ranges, antitank rocket range impact areas, demolition ranges, 40-mm rifle grenade ranges, hand grenade ranges, and firing points for the various weapons. The chemical and physical characteristics of energetic residues at these ranges differ substantially. For example, at antitank rocket ranges, nitroglycerin (NG) is deposited near the firing point and is present at the highest concentrations behind the firing line due to back blast. The energetic residue present at the highest concentrations in the impact area of this type of range is mainly 1,3,5,7-octahydro-1,3,5,7-tetranitrotetrazocine (HMX) from physically breached casings of the antitank rocket's warhead (Jenkins et al. 1999). For artillery range impact areas, however, the major residues are either 2,4,6-trinitrotoluene (TNT) or Composition B (60% 1,3,5-hexahydro-1,3,5-trinitrotriazine [RDX], 39 % TNT) from the warheads of artillery or mortar rounds. The major residues at artillery ranges reside as distributed sources associated with rounds that have undergone low-order (partial) detonation (Pennington et al. 2002, 2003).

A study was conducted at Canadian Force Base–Valcartier (CFB–Valcartier) in which various sampling protocols were evaluated with regard to their ability to provide representative samples that would provide estimates of mean concentration (Jenkins et al. 2004). Both discrete and multi-increment composite samples were collected within 10-m \times 10-m areas at the firing point and impact area where training is conducted with antitank rockets. The largest sampling problem to be overcome was that residues were distributed heterogeneously over the area, resulting in a range of concentrations of greater than two orders of magnitude in discrete samples collected at both the firing point (NG) and impact area (HMX). From these results it is clear that collection of discrete samples to estimate a mean concentration for areas as small as 100 square meters is very unreliable and the individual values will underestimate the mass of residue loading in most cases.

2 OBJECTIVES

The major objective of this work was to thoroughly characterize an area within an artillery range where the major contamination source was a low-order detonation of a mortar or artillery round to understand the distribution of residues of high explosives. Specifically an attempt was made to estimate the short-range heterogeneity in analyte concentrations and investigate various alternatives to obtain samples representative of the mean analyte concentrations within this type of area. Previous work has shown that low-orders are an anomaly even in high-use target areas. Therefore, in general, the concentrations of energetic residues are anticipated to be very low (Jenkins et al. 1998, 2001; Pennington et al. 2001, 2002, 2003). However, when rounds low-order, a hot spot of residue concentrations is often created, thereby presenting a unique sampling challenge.

A second objective was to collect surface soil samples spatially around an artillery target to determine whether residue concentrations were distributed in a similar manner as found for targets at antitank ranges. For antitank ranges, sampling studies at a number of installations have shown that there is a strong concentration gradient for HMX with the highest concentrations next to the target (Jenkins et al. 1997, 1998; Pennington et al. 2002; Thiboutot et al. in press).

3 SOIL SAMPLE COLLECTION AND ANALYSIS

Soil sample collection in an area near a low-order detonation of an 81-mm mortar

This study was conducted at an artillery impact range at Fort Polk, Louisiana, from 25 June to 29 June 2003. Initially all field personnel accompanied the AEC/CHPPM sampling team as they traversed the impact range on foot to collect soil samples at a number of predetermined locations according to their stratified random sampling strategy. At an access road that was downhill from an artillery target, small chunks of what appeared to be explosives residue were observed on the soil surface (Fig. 1). Using the EXPRAY kit (a field test kit designed to detect and classify energetic residues [Plexus Scientific, Silver Spring, Maryland]), these chunks were shown to contain both a nitroaromatic (probably TNT) and a nitramine/nitrate ester (probably RDX) in the chunks of residue found on the ground. Subsequent laboratory analysis at CRREL confirmed the proper ratio of RDX/TNT for Composition B. After inspecting the surface of the soil in the vicinity of these chunks, additional pieces of explosives material were observed. The presence of a fin and casing fragment indicated that the explosives residues originated from an 81-mm mortar that had undergone a low-order detonation.



Figure 1. Chunks of Composition B from the partial detonation of an 81-mm mortar round found in the artillery impact area at Fort Polk, Louisiana.



Figure 2. Sampling a 10-m \times 10-m grid area encompassing pieces of chunk explosive from a low-order 81-mm mortar round. The tank target is uphill and in the background.

Because we were interested in studying the distribution of explosives residues near low-order detonations, we located a 10-m \times 10-m sampling grid along the road, encompassing the pieces of chunk explosive that we had visually identified (Fig. 2). The center of this grid was about 30 m downslope from a heavily used tank target. A diagram of the area is shown in Figure 3. From the surface topography we do not believe that any surface runoff from the tank target located upslope and left of this grid (as seen from the road) would pass over this grid.

The 10-m \times 10-m sampling grid was further subdivided into 100 1-m \times 1-m minigrids (Fig. 4). Within each minigrid a discrete sample was collected from the surface and stored in a 4-oz amber glass jar. These samples and all subsequent samples were obtained using a coring device (Fig. 5) developed at CRREL (Walsh 2004). All core sample increments were of the top 2.5 cm and were 4.8 cm in diameter. If visible pieces of explosive were present in a given minigrid, the material was gathered and weighed with a portable balance (Fig. 6), and we collected the soil sample adjacent to where the largest piece of solid explosive had been located. In minigrids that did not have visible solid explosive, samples were collected at a randomly selected location within the 1-m \times 1-m area. Duplicate discrete soil samples were collected adjacent to the initial discrete sampling location in 20 randomly selected minigrids. Within these same 20 randomly selected minigrids, a ten-increment composite was also collected at

randomly selected locations and stored in a 32-oz glass jar. The piece of the low-order 81-mm mortar (tail fin and casing fragment) that was observed was located at the boundary of minigrids 3 and 13.

After all of the visible explosive was removed, and the minigrid discrete and ten-increment composite samples were collected, ten 25-increment composite samples were collected from the entire 10-m \times 10-m grid using a random sampling strategy (moving in random directions after the collection of each increment, i.e., analogous to random number generator). Five different individuals collected these samples using the soil corers described above. Samples were stored in 64-oz glass jars.

Beyond the boundaries of the 10-m \times 10-m grid, ten-increment composite samples were collected from lines parallel to and outward from the four sides of the grid at distances of 2 m, 5 m, and 10 m as shown in Figure 4. These composite samples were stored in 32-oz glass bottles.

Sampling near a tank target uphill from the 10-m \times 10-m grid

A heavily impacted artillery target was located about 30 m uphill and to the left (as seen from the road) of the 10-m \times 10-m sampling grid. Twelve sampling grids were established as shown in Figure 3. The boundaries of the four inside grids were from the perimeter of the tank target to 2 m from the edge. The eight outer grids were located from 2 m to 5 m from the edge of the target. A set of ten-increment surface soil composite samples was collected from within grid boundaries using a random sampling strategy similar to that described above. These composite samples were stored in 32-oz glass bottles.

Soil sample analysis

Soil samples were returned to CRREL and air-dried at room temperature. The discrete and composite samples were processed differently because the sample masses were different.

Discrete samples were dried in 4-oz amber containers, weighed, passed through a #10 (2-mm) sieve to remove oversize material, the sieved portion weighed, and returned to the 4-oz containers. Discrete samples were not subsampled, rather the entire sample was extracted as follows. A volume of acetonitrile in milliliters (mL), approximately double the mass of the sample in grams, was added to each 4-oz jar unless the sample was too large (greater than 60 g) (Hewitt and Walsh 2003). For those cases the sample was transferred to an 8-oz jar and acetonitrile was added. All jars were capped and placed on a tabletop shaker overnight (18 hours @ 150 rpm). The samples were removed from the

shaker and allowed to settle for at least an hour. An aliquot of each extract was filtered through a 0.45- μm Millex FH filter and placed in a 7-mL amber glass vial. Vials were stored in a refrigerator until analyzed.

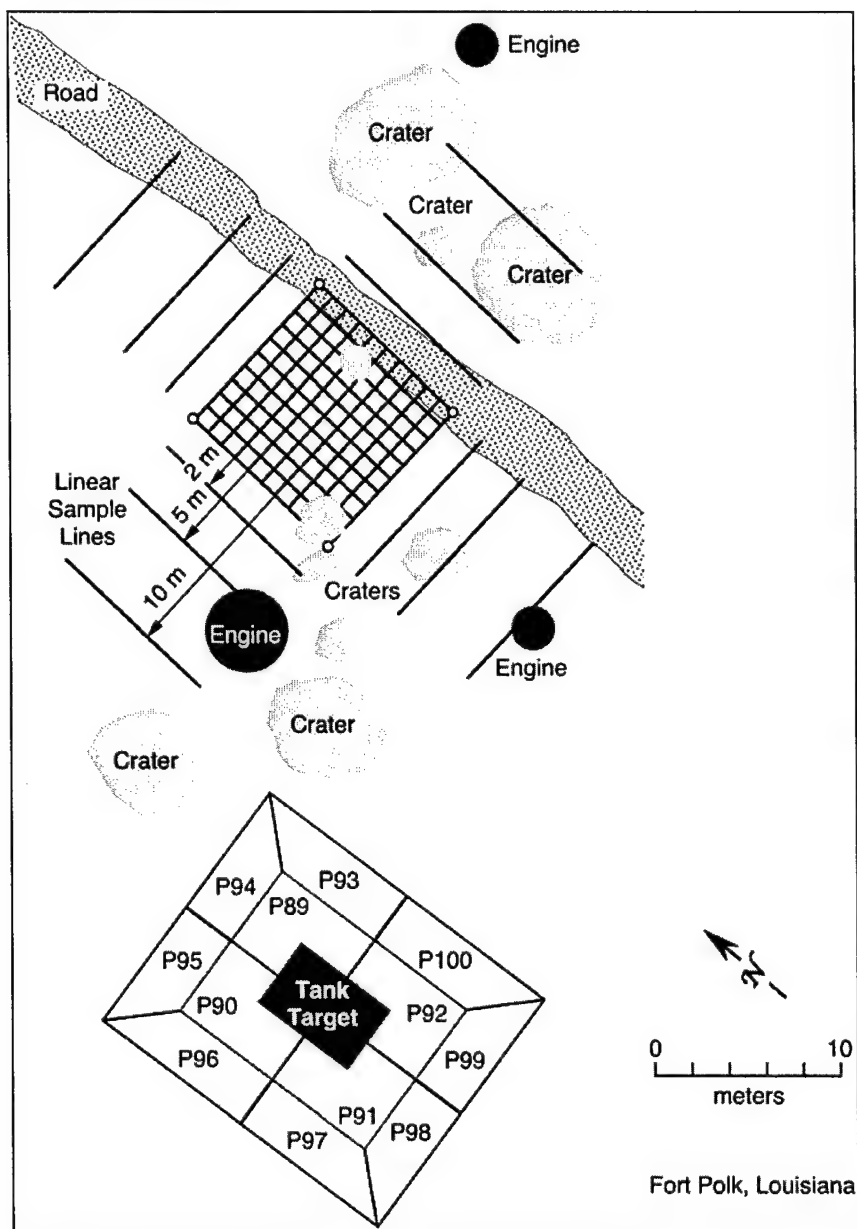


Figure 3. Relationship of a 10-m x 10-m sampling grid and a tank target in the artillery impact area at Fort Polk, Louisiana.

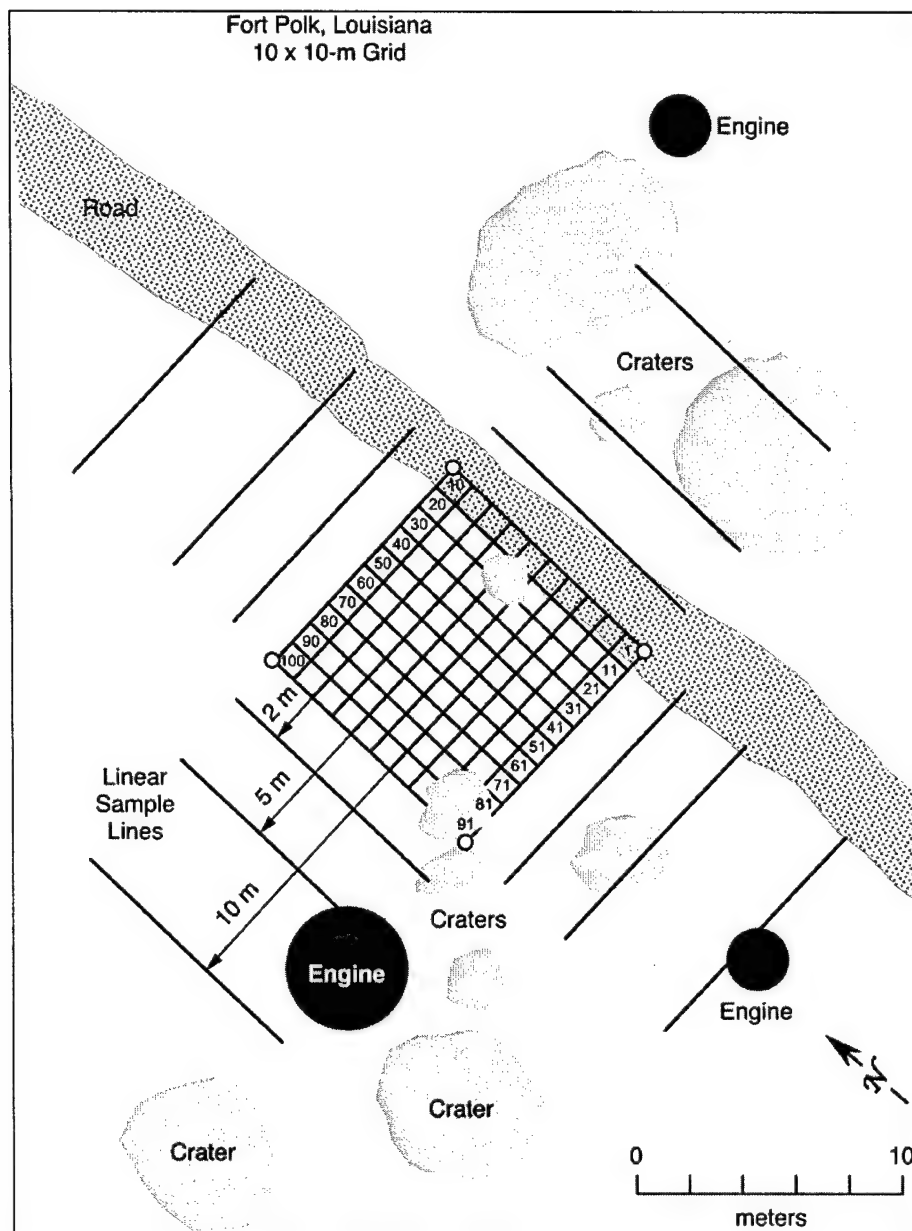


Figure 4. 10-m × 10-m sampling grid subdivided into 100-m² minigrids and linear sampling lines around the major grid area.



Figure 5. Coring device used to collect soil samples at Fort Polk, Louisiana.



Figure 6. Collecting and weighing visible pieces of Composition B found around a partial detonation of an 81-mm mortar.

Composite soil samples were placed on sheets of aluminum foil to air-dry. Dried samples were weighed and sieved through a #10 sieve. The material that passed the sieve was weighed and ground in a Lab TechEssa LM2 (LabTech Essa Pty. Ltd., Bassendean, WA, Australia) puck mill grinder for 60 seconds. After grinding, composite samples were mixed thoroughly and then spread out to form a 1- to 2-cm-thick layer. A subsample then was obtained by collecting at least 30 increments randomly from the ground material for a mass of about 10 g. For every tenth sample, an additional subsample was collected in an identical manner to enable an assessment of subsampling uncertainty. Each 10-g subsample was extracted with 20 mL of acetonitrile in an ultrasonic bath overnight at room temperature. After sonication, samples were removed from the bath and allowed to settle for at least an hour. An aliquot was then removed, filtered, and placed in a 7-mL amber vial for storage in a refrigerator.

Commercial sand was used as a laboratory processing blank. This blank soil was ground, subsampled, and extracted with each batch (i.e., approximately 20 samples) of field samples. A standard soil obtained from the U.S. Army Environmental Center was used for preparation of the laboratory control sample. This soil was spiked with a suite of target analytes and was used to assess recovery.

The extracts from both the discrete and composite samples were all analyzed using the general procedures of SW 846 Method 8330 (EPA 1994). For this analysis, an aliquot of each sample was diluted one part extract to three parts reagent-grade water. Analysis was conducted on a modular RP-HPLC system from Thermo Finnigan composed of a SpectraSYSTEM Model P1000 isocratic pump, a SpectraSYSTEM UV2000 dual wavelength UV/VS absorbance detector set at 210 and 254 nm (cell path 1 cm), and a SpectraSYSTEM AS300 auto-sampler. Samples were introduced by overfilling a 100- μ L sampling loop. Separations were made on a 15-cm \times 3.9-mm (4- μ m) NovaPak C-8 column (Waters Chromatography Division, Milford, Massachusetts) maintained at 28°C and eluted with 15:85 isopropanol/water (v/v) at 1.4 mL/min. Concentrations were estimated from peak heights compared to commercial multianalyte standards (Restek). If concentrations exceeded 20 ppm, an aliquot of the original extract was diluted appropriately with additional acetonitrile prior to the 1 to 4 dilution with reagent-grade water. Estimates of detection limits for the target analytes for this method are given in Table 1.

For low-concentration (< 0.2 mg/kg) samples, a second analysis was conducted by GC-ECD following the general procedure outlined in SW846 Method 8095 (EPA 1999). These analyses were conducted on an HP 6890 Gas Chromatograph equipped with a micro ECD detector. Direct injection of 1 μ L of soil extract was made into a purged packed inlet port (250°C) equipped with a

deactivated Restek Uniliner. Primary separation was conducted on a 6-m- × 0.53-mm-ID fused-silica column, with a 1.5- μ m film thickness of 5% (phenyl)-methylsiloxane (Rtx-5 from Restek, Bellefonte, Pennsylvania).

The GC oven was temperature-programmed as follows: 100°C for 2 min, 10°C/min ramp to 280°C. The carrier gas was hydrogen at 10 mL/min (linear velocity approximately 90 cm/sec). The ECD detector temperature was 310°C and the makeup gas was nitrogen flowing at 45 mL/min. If a peak was observed in the retention window for a specific signature compound, the extract was reanalyzed on a confirmation column, 6-m × 0.53-mm ID having a 1.5- μ m film thickness of a proprietary polymer (Rtx-TNT-2 from Restek). The GC oven was temperature-programmed as follows: 130°C for 1 min, 10°C/min ramp to 280°C. The carrier gas was helium at 20 mL/min (linear velocity approximately 180 cm/sec) and the nitrogen makeup gas was flowing at 60 mL/min. Inlet and detector temperature were the same as above. Multianalyte standards were purchased from Restek and the instrument was calibrated over five concentrations. Estimates of the detection limits for the GC-ECD method are given in Table 1.

Table 1. Explosives detection limits for soil and water.

Analyte	Soil (μ g/kg)		Water (μ g/L)
	RP-HPLC	GC-ECD	GC-ECD
HMX	26	10	0.047
RDX	34	6	0.035
TNB	16	3	0.016
TNT	16	2	0.017
2,6-DNT	19	2	0.009
2,4-DNT	28	2	0.010
2ADNT	38	2	0.028
4ADNT	32	2	0.018
NG	20	10	0.20
DNA	Co-elutes with NB	2	0.019
DNB	15	2	0.010
Tetryl	100	10	0.025
PETN	56	16	0.24

4 RESULTS

Quality control

Results for the analysis of laboratory duplicate soils samples, blank soils, and laboratory control (spike) samples (LCS) are shown in Table 2. No target analytes were detected in any of the blank soils. The recovery for the spiked LCS samples ranged from 83% to 108%, with a mean value of 97.6%. Eight composite samples had laboratory duplicates removed for analysis. In most cases the results for the laboratory duplicates indicated that the results were very reproducible. For soil sample P-58, however, the agreement of the duplicates was poor, so we examined the sample further. Upon careful inspection it was obvious that the soil sample was not adequately ground because the texture was not uniform. We reground this sample and duplicate subsamples were taken, extracted, and analyzed. The original data for sample P-58 and data for the sample after regrounding are given in Table 2.

Table 2. Quality control soil samples for Fort Polk study. (Analysis by RP-HPLC and GC-ECD [shaded].)

Lab #	Soil concentration (mg/kg)								
	HMX	TNB	RDX	TNT	NG	2,4-DNT	2,6-DNT	4ADNT	2ADNT
Laboratory duplicate samples									
P-58	0.162	<d	0.432	0.032	0.102	<d	0.086	0.086	0.050
P-58LD	<d	<d	0.118	0.036	1.63	0.176	0.008	0.010	0.007
After regrounding P-58									
P-58-1-1	0.040	<d	0.116	0.034	0.05	0.372	0.018	0.006	0.004
P-58-1-2	0.050	<d	0.138	0.032	0.20	0.376	0.006	0.006	0.008
P-72	<d	<d	0.062	<d	<d	<d	<d	<d	<d
P-72LD	<d	<d	0.060	<d	<d	<d	<d	0.002	0.002
P-80	0.614	<d	4.56	1.20	<d	<d	<d	0.158	0.156
P-80LD	0.614	<d	4.58	1.20	<d	<d	<d	0.140	0.156
P-89	15.1	<d	16.4	1.21	<d	<d	<d	0.246	0.328
P-89LD	15.4	<d	15.4	1.17	<d	<d	<d	0.244	0.288
P-98	0.224	<d	1.23	2.24	<d	<d	<d	0.612	0.890
P-98LD	0.226	<d	1.22	2.20	<d	<d	<d	0.610	0.860
D-39C	0.064	<d	0.512	0.008	<d	<d	<d	0.040	0.048
D-39C LD	0.062	<d	0.532	0.010	<d	<d	<d	0.038	0.040

Table 2 (cont'd).									
Lab #	Soil concentration (mg/kg)								
	HMX	TNB	RDX	TNT	NG	2,4-DNT	2,6-DNT	4ADNT	2ADNT
D-82C	0.674	<d	4.46	0.462	<d	<d	<d	0.306	0.316
D-82C LD	0.714	<d	4.54	0.474	<d	<d	<d	0.344	0.338
G-1-6	8.58	<d	80.0	21.2	<d	<d	<d	0.808	0.798
G-1-6 LD	7.96	<d	75.8	20.4	<d	<d	<d	0.728	0.784
Lab blanks									
LB-1	<d	<d	<d	<d	<d	<d	<d	<d	<d
LB-2	<d	<d	<d	<d	<d	<d	<d	<d	<d
LB-3	<d	<d	<d	<d	<d	<d	<d	<d	<d
LB-4	<d	<d	<d	<d	<d	<d	<d	<d	<d
LB-5	<d	<d	<d	<d	<d	<d	<d	<d	<d
LB-6	<d	<d	<d	<d	<d	<d	<d	<d	<d
Spike samples									
LCS-1	0.542	0.548	0.505	0.445	—	0.509	0.445	0.539	0.534
LCS-2	0.488	0.488	0.443	0.415	—	0.475	0.477	0.482	0.463
Percent recoveries									
LCS-1	108%	110%	101%	89%	—	102%	89%	108%	107%
LCS-2*	98%	98%	89%	83%	—	95.0%	95.4%	96.4%	92.6%
* LCS-2 was analyzed with a different lot of samples and calibration than LCS-1.									

After regrinding, the agreement of the data for all analytes is much improved, confirming that the initial problem with this soil was due to inadequate grinding, most likely because initially the grinding bowl contained too much soil. Subsequently, the laboratory protocol was changed to grinding aliquots of no more than 500 g (the manufacturer-recommended cutoff is 800 g).

Because we were concerned that other soil samples from Fort Polk also may have suffered this problem, we inspected all the composite samples and found five that also appeared to be inadequately ground. These samples also were reground and triplicate subsamples were analyzed. The original results for these samples and the results for the triplicate reground samples are given in Table 3. For these samples, changes in the analytical results were much smaller and often insignificant (i.e., < 15% RPD).

Table 3. Analytical results for Fort Polk samples that were analyzed, reground, and subsamples analyzed in triplicate.

Lab #	Soil concentration (mg/kg)				
	HMX	RDX	TNT	4ADNT	2ADNT
D-14C regrinding	18.5	139	15.5	2.24	2.88
1	17.7	127	14.7	2.20	2.66
2	17.7	128	14.9	2.14	2.72
3	17.5	126	14.4	2.08	2.56
D-46C regrinding	0.294	1.85	0.158	0.098	0.124
1	0.320	1.94	0.242	0.070	0.142
2	0.316	1.91	0.206	0.064	0.122
3	0.308	1.86	0.198	0.084	0.114
D-51C regrinding	3.04	21.6	2.68	0.640	0.460
1	3.20	22.6	2.78	0.606	0.648
2	3.24	22.4	2.80	0.562	0.658
3	3.26	22.8	2.70	0.588	0.636
D-82C	0.674	4.46	0.462	0.306	0.316
D-82C LD regrinding	0.714	4.54	0.474	0.344	0.338
1	0.666	4.18	0.494	0.276	0.266
2	0.656	4.20	0.486	0.270	0.258
3	0.678	4.24	0.506	0.278	0.258
D-87C regrinding	0.228	1.504	37.4	0.278	0.260
1	0.220	1.468	35.2	0.260	0.230
2	0.218	1.422	34.2	0.272	0.228
3	0.240	1.428	34.2	0.264	0.226

Grid samples from the area near a low-order 81-mm mortar detonation

Analytical results for the 100 discrete minigrid samples collected from within a 10-m × 10-m grid from the area where an 81-mm mortar round had low-ordered are presented in Table 4. RDX, HMX, TNT, 2ADNT, and 4ADNT were detected in nearly all of the discrete surface soil samples. RDX was present at the highest concentration, with surface soil concentrations ranging over almost five orders of magnitude from 0.037 to 2,390 mg/kg. The median RDX concentration

was 1.79 mg/kg, but because there were several very high concentrations, mean concentration was 70.9 mg/kg. HMX concentrations in these discrete minigrid samples ranged from less than detection limits (0.01 mg/kg) to 253 mg/kg. The median ratio of HMX to RDX was 0.176, which is slightly higher than expected (about 0.11) for HMX as an impurity in RDX. This ratio suggests that weathering has resulted in the preferential dissolution of the more soluble RDX.

Table 4. Results from the analysis of discrete samples from 100 1-m × 1-m minigrids in an area near a low-order 81-mm mortar detonation at the impact range at Fort Polk. (Analysis by RP-HPLC and GC-ECD [shaded].)

Field #	Comp. B found (g)	Soil concentration (mg/kg)					Ratio		
		HMX	RDX	TNT	4ADNT	2ADNT	TNT/RDX	HMX/RDX	2ADNT/4ADNT
D1		0.893	8.86	2.72	0.124	0.170	0.307	0.101	1.38
D2	0.1	0.639	3.50	0.071	0.178	0.198	0.020	0.182	1.11
D3	50.2	1.12	5.02	0.072	0.563	0.590	0.014	0.223	1.05
D4	0.1	7.50	42.7	6.53	0.418	0.410	0.153	0.176	0.98
D5	0.1	44.4	385	147	3.18	3.10	0.382	0.115	0.97
D6	0.3	2.46	24.9	0.095	0.707	0.715	0.004	0.099	1.01
D7		0.740	3.64	0.064	0.115	0.125	0.018	0.203	1.10
D8		0.190	0.965	0.003	0.047	0.068	0.004	0.197	1.44
D9		0.126	0.526	0.002	0.028	0.029	0.004	0.239	1.06
D10		0.041	0.161	0.004	0.009	0.008	0.026	0.252	0.89
D11		0.230	1.18	0.003	0.099	0.129	0.003	0.195	1.30
D12		0.235	1.03	0.023	0.510	0.633	0.023	0.227	1.24
D13	28.7	11.3	64.3	13.4	1.32	1.75	0.208	0.176	1.33
D14	7.8	60.2	557	164	3.30	3.61	0.294	0.108	1.09
D15	5.5	189	1790	489	16.3	15.3	0.273	0.106	0.94
D16	17	253	2390	1560	0.090	0.125	0.653	0.106	1.38
D17	0.3	1.27	11.3	2.50	0.392	0.344	0.222	0.113	0.88
D18		0.371	1.65	0.084	0.265	0.244	0.051	0.224	0.92
D19		0.075	0.335	0.024	0.036	0.042	0.070	0.225	1.16
D20		0.028	0.263	0.001*	0.009	0.011	0.004	0.108	1.20
D21	1.4	5.74	48.3	29.7	0.349	0.476	0.615	0.119	1.36
D22	0.2	1.42	13.3	4.46	0.366	0.457	0.334	0.106	1.25
D23		0.484	3.36	0.901	0.622	0.706	0.268	0.144	1.13
D24	3	0.925	6.93	0.240	1.50	1.64	0.035	0.133	1.10
D25	48.2	97.4	889	321	5.90	5.47	0.362	0.110	0.93
D26	13.1	3.56	21.8	0.932	1.76	1.91	0.043	0.164	1.09

Table 4 (cont'd). Results from the analysis of discrete samples from 100 1-m × 1-m minigrids in an area near a low-order 81-mm mortar detonation at the impact range at Fort Polk. (Analysis by RP-HPLC and GC-ECD [shaded].)

Field #	Comp. B found (g)	Soil concentration (mg/kg)					Ratio		
		HMX	RDX	TNT	4ADNT	2ADNT	TNT/RDX	HMX/RDX	2ADNT/4ADNT
D27	1	0.528	3.75	0.022	0.153	0.182	0.006	0.141	1.19
D28	0.4	0.185	0.618	0.002	0.038	0.045	0.003	0.299	1.17
D29		0.077	0.193	0.004	0.034	0.040	0.022	0.402	1.19
D30		0.016	0.081	0.001*	0.008	0.009	0.012	0.198	1.13
D31	0.4	0.419	1.65	0.003	0.195	0.253	0.002	0.255	1.30
D32		0.561	1.56	0.017	0.394	0.456	0.011	0.358	1.16
D33		0.476	8.51	0.247	0.710	0.636	0.029	0.056	0.89
D34		1.42	10.6	3.47	0.566	0.691	0.329	0.135	1.22
D35		0.215	2.24	0.106	0.878	1.03	0.047	0.096	1.17
D36		3.35	25.2	6.88	1.04	1.13	0.273	0.133	1.08
D37		2.04	7.15	0.168	0.590	0.698	0.023	0.285	1.18
D38		0.082	0.248	0.003	0.038	0.051	0.014	0.332	1.36
D39		0.067	0.175	0.009	0.049	0.053	0.049	0.385	1.08
D40		0.005*	0.037	0.001*	0.009	0.013	0.027	0.135	1.50
D41		2.17	7.52	0.871	0.848	0.910	0.116	0.289	1.07
D42		1.66	5.65	0.132	0.586	0.777	0.023	0.294	1.33
D43		0.304	1.97	0.027	0.342	0.348	0.014	0.154	1.02
D44	0.1	0.108	0.571	0.009	0.347	0.471	0.016	0.189	1.36
D45	0.1	1.76	4.84	0.062	0.461	0.510	0.013	0.365	1.11
D46		2.52	19.9	3.94	0.711	0.668	0.198	0.126	0.94
D47		0.232	0.825	0.014	0.098	0.134	0.017	0.281	1.36
D48		0.036	0.122	0.004	0.020	0.027	0.031	0.296	1.38
D49		0.092	1.46	0.004	0.015	0.018	0.002	0.063	1.18
D50		0.020	0.070	0.001*	0.010	0.010	0.014	0.292	1.00
D51	0.2	34.9	331	81.6	2.23	2.15	0.246	0.105	0.96
D52		1.88	9.70	0.189	0.350	0.488	0.019	0.194	1.39
D53	16.3	1.54	3.96	0.434	0.915	1.17	0.110	0.390	1.28
D54		0.725	1.44	0.014	0.253	0.246	0.010	0.504	0.97
D55		0.517	3.67	1.31	0.108	0.104	0.356	0.141	0.96
D56		0.043	0.243	0.001	0.023	0.027	0.004	0.176	1.17
D57		0.669	3.21	0.005	0.037	0.068	0.002	0.208	1.82
D58		0.094	0.254	0.006	0.035	0.054	0.023	0.369	1.53
D59		0.086	1.03	0.012	0.028	0.027	0.011	0.083	0.95
D60		0.005*	0.073	0.001*	0.012	0.016	0.014	0.069	1.38

Table 4 (cont'd).

Field #	Comp. B found (g)	Soil concentration (mg/kg)					Ratio		
		HMX	RDX	TNT	4ADNT	2ADNT	TNT/RDX	HMX/RDX	2ADNT/ 4ADNT
D61		2.55	12.7	0.924	1.41	1.76	0.073	0.201	1.25
D62	1.1	16.5	138	76.1	3.14	3.76	0.552	0.120	1.20
D63	0.9	6.38	53.7	28.2	0.919	1.03	0.525	0.119	1.12
D64	0.5	0.736	3.85	0.028	0.893	1.15	0.007	0.191	1.29
D65	1.2	0.585	4.94	0.656	0.156	0.176	0.133	0.119	1.13
D66	0.1	0.269	1.22	0.007	0.130	0.168	0.005	0.221	1.29
D67		0.535	4.63	1.70	0.063	0.079	0.368	0.115	1.25
D68		0.117	0.470	0.003	0.013	0.022	0.006	0.248	1.70
D69		0.188	2.41	0.007	0.023	0.025	0.003	0.078	1.07
D70		0.216	1.06	0.044	0.063	0.173	0.041	0.203	2.75
D71		5.91	30.8	5.41	0.486	0.515	0.176	0.192	1.06
D72		0.468	1.40	0.054	0.755	0.813	0.039	0.333	1.08
D73		2.23	12.5	1.32	0.619	0.736	0.106	0.179	1.19
D74		0.044	0.342	0.006	0.117	0.173	0.016	0.128	1.48
D75		0.005*	0.074	0.002	0.019	0.025	0.021	0.067	1.33
D76		0.237	1.11	0.007	0.047	0.086	0.007	0.212	1.84
D77		0.109	0.180	0.001*	0.016	0.022	0.006	0.605	1.36
D78		0.022	0.076	0.001*	0.015	0.018	0.013	0.286	1.25
D79		0.667	7.11	2.58	0.087	0.076	0.363	0.094	0.87
D80		0.058	0.187	0.004	0.017	0.031	0.023	0.311	1.83
D81		0.089	0.805	0.045	0.238	0.235	0.056	0.110	0.99
D82		2.80	24.1	3.29	1.11	1.09	0.136	0.116	0.98
D83		1.55	7.73	0.783	0.479	0.550	0.101	0.201	1.15
D84		0.098	0.539	0.003	0.102	0.129	0.005	0.182	1.26
D85		0.095	0.260	0.003	0.045	0.045	0.010	0.364	1.00
D86		0.031	0.233	0.001*	0.018	0.021	0.004	0.134	1.15
D87		0.076	0.366	0.003	0.021	0.049	0.009	0.208	2.38
D88		0.225	1.93	0.396	0.075	0.069	0.205	0.116	0.92
D89		0.083	0.731	0.002	0.012	0.026	0.003	0.114	2.17
D90		0.108	0.138	0.004	0.022	0.022	0.031	0.781	1.00
D91		1.57	17.1	1.68	0.739	0.769	0.098	0.092	1.04
D92		0.358	1.27	0.093	0.095	0.126	0.073	0.281	1.32
D93		0.065	0.829	0.038	0.154	0.157	0.046	0.078	1.02
D94		0.111	0.908	0.232	0.031	0.052	0.256	0.122	1.68
D95		0.777	10.9	0.015	0.103	0.142	0.001	0.071	1.38
D96		0.516	4.44	1.86	0.054	0.109	0.419	0.116	2.00

Table 4 (cont'd). Results from the analysis of discrete samples from 100 1-m × 1-m minigrids in an area near a low-order 81-mm mortar detonation at the impact range at Fort Polk. (Analysis by RP-HPLC and GC-ECD [shaded].)

Field #	Comp. B found (g)	Soil concentration (mg/kg)					Ratio		
		HMX	RDX	TNT	4ADNT	2ADNT	TNT/RDX	HMX/RDX	2ADNT/4ADNT
D97		0.059	0.437	0.102	0.019	0.034	0.232	0.135	1.81
D98		0.047	0.354	0.003	0.024	0.038	0.009	0.134	1.60
D99		0.170	1.52	0.001*	0.016	0.020	0.001	0.112	1.23
D100		0.005*	0.067	0.001*	0.013	0.022	0.015	0.074	1.60
Max		253	2390	1560	16.3	15.3	0.653	0.781	2.75
Min		0.005	0.037	0.001	0.008	0.008	0.001	0.056	0.873
Median		0.395	1.79	0.044	0.120	0.169	0.025	0.176	1.18
Mean		7.89	70.9	29.7	0.626	0.663	0.107	0.195	1.25

* Concentration was nondetectable; one-half the detection limit was used for mathematical computation.

Note: Because the distributions depart greatly from normal as shown by the large discrepancy between median and mean and by the histogram, it is not possible to compute useful standard deviations.

TNT concentrations in these samples were always lower than RDX and ranged from less than 0.002 mg/kg to 1,560 mg/kg (Table 4). The mean ratio of TNT to RDX was 0.107. For non-weathered Composition B, the ratio should be about 0.7, indicating that the TNT present in these samples has been subject to preferential dissolution and environmental transformation. Even so, based on our experience, the presence of TNT in these samples is consistent with the contamination source being Composition B.

The two most common environmental transformation products of TNT, 2ADNT and 4ADNT, were detected in all 100 minigrid samples, even in samples where the TNT concentration was below detection limits. In fact, the median concentrations of these two transformation products were actually higher than TNT for these 100 samples. The median ratio of 2ADNT/4ADNT was 1.18 and, from our experience, this ratio is typical for these compounds in surface soil samples.

The distribution of the 100 discrete minigrid concentrations for RDX is presented as a histogram in Figure 7; the distribution is clearly non-Gaussian. A histogram presenting the log of the concentration versus frequency is shown in Figure 8.

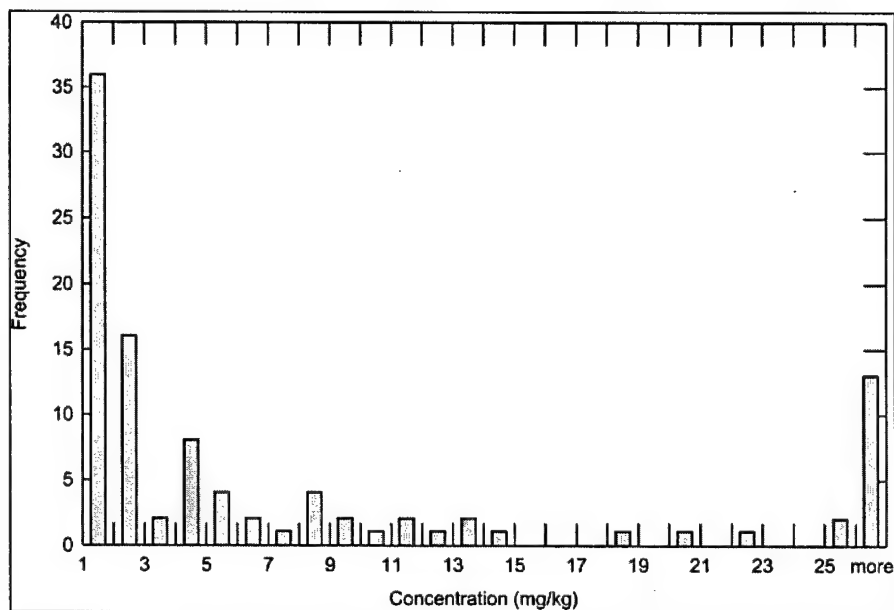


Figure 7. Distribution of soil RDX concentrations from 100 discrete soil samples taken in the 1-m \times 1-m minigrids.

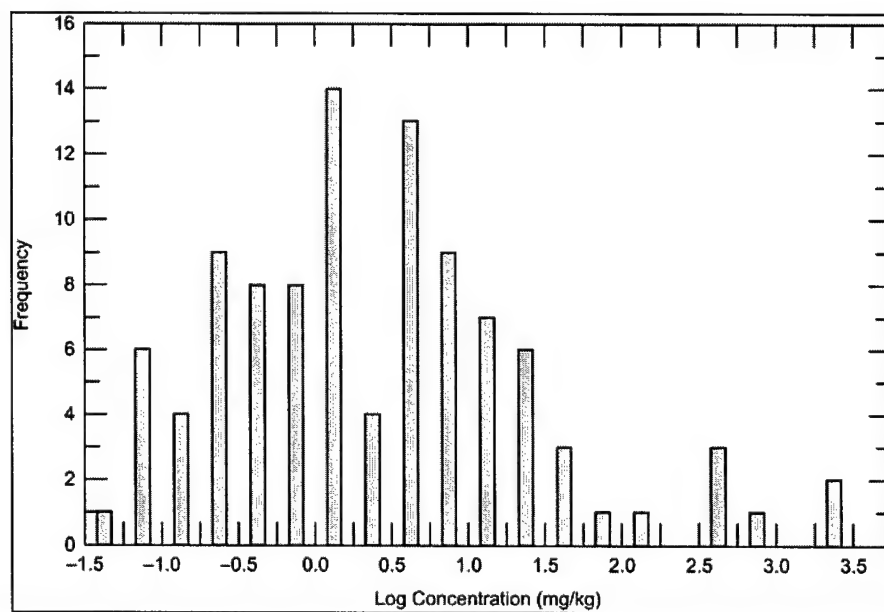


Figure 8. Distribution of the log soil RDX concentrations from 100 discrete soil samples taken from 1-m \times 1-m minigrids.

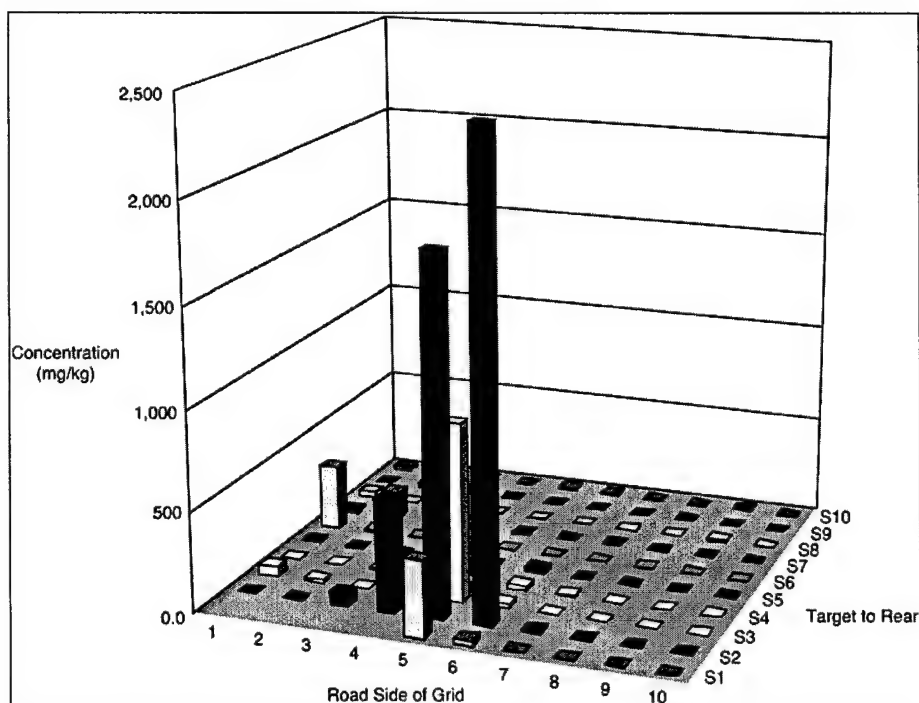


Figure 9. Soil concentration of RDX relative to its location within the 10-m \times 10-m grid.

A plot of the RDX concentrations in surface soil versus position within the 10-m \times 10-m grid is presented in Figure 9. It appears from these data that one major hot spot within this grid is centered at minigrid D15 and a smaller one near the edge of the grid is centered at grid D62. These apparent hot spots can also be seen in the weights of Composition B recovered from the individual minigrids (Table 4), although the centers of the hot spots do not agree exactly (Figure 10). These hot spots coincide with the high concentration population in Figure 8 and are spatially distinct. The lack of complete agreement shows that the presence of visible residues on the surface is not necessarily a prerequisite to finding high concentrations in the soil, i.e., there can be areas adjacent to visible residues with high concentrations of fine non-visible particles.

Mass of Composition B (g) Collected in Grid										RDX Concentration in Surface Soil (mg/kg)									
										17.1	1.27	0.829	0.908	10.9	4.44	0.437	0.354	1.52	0.067
										0.805	24.1	7.73	0.539	0.260	0.233	0.366	1.93	0.731	0.138
										30.8	1.40	12.5	0.342	0.074	1.11	0.18	0.076	7.11	0.187
										12.7	138	53.7	3.85	4.94	1.22	4.63	0.470	2.41	1.06
										331	9.70	3.96	1.44	3.67	0.243	3.21	0.254	1.03	0.073
										7.52	5.65	1.97	0.571	4.84	19.9	0.825	0.122	1.46	0.070
										1.65	1.56	8.51	10.6	2.24	25.2	7.15	0.248	0.175	0.037
										48.3	13.3	3.36	6.93	889	21.8	3.75	0.618	0.193	0.081
										1.18	1.03	64.3	557	1790	2390	11.3	1.65	0.335	0.263
										8.86	3.50	5.02	42.7	385	24.9	3.64	0.96	0.526	0.161
1	2	3	4	5	6	7	8	9	10	1	2	3	4	5	6	7	8	9	10

Fort Polk, Louisiana
Sampling Grid (10 m x 10 m)

Figure 10. Weight of Composition B and soil RDX concentration and their relative position in the sampling grid.

Comparison of field duplicate discrete and ten-increment composites for minigrid samples

Results for the duplicate discrete samples from the 20 randomly selected minigrids are presented in Table 5. An inspection of these data indicates that agreement between replicates is analyte-dependant. For example, while there is greater than a two-orders-of-magnitude difference between the RDX, TNT, and HMX concentrations for the field duplicates taken from minigrids 80 and 100, the discrepancy between 4ADNT and 2ADNT is less than a factor of three. This anomaly can be explained by the physical state of these analytes. RDX, TNT, and HMX are present as crystalline particulates, whereas 4ADNT and 2ADNT are formed only following dissolution and subsequent biotransformation. Therefore, one group of energetic materials exists as discrete particles while the other exists on the surfaces of soil grains and has had the opportunity to disperse into the surrounding substrate. Because of this phenomenon, individual cell discrete data are inadequate to represent areas as small as one square meter when the analytes remain in a crystalline particulate state. Thus, any type of site characterization based on discrete samples for RDX, TNT, and HMX would not be valid.

Table 5. Results from the analysis of duplicate discrete and ten-increment composite samples from 20 randomly chosen minigrids near location of low-order mortar detonation. (AcN extraction with RP-HPLC and GC-ECD [shaded] analysis.)

Field #	Soil concentration (mg/kg)				
	HMX	RDX	TNT	4ADNT	2ADNT
D14*	60.2	557	164	3.30	3.61
D14-FD†	23.5	167	3.57	2.43	2.94
C14**	18.5	139	15.5	2.24	2.88
D19	0.075	0.335	0.024	0.036	0.042
D19-FD	0.067	0.225	<d	0.046	0.061
C19	0.364	2.40	0.196	0.106	0.110
D27	0.528	3.75	0.022	0.153	0.182
D27-FD	0.342	1.68	0.025	0.240	0.278
C27	3.24	24.2	6.10	0.562	0.604
D28	0.185	0.618	0.002	0.038	0.045
D28-FD	0.195	0.693	<d	0.058	0.080
C28	3.24	26.8	7.46	0.732	0.712
D29	0.077	0.193	0.004	0.034	0.040
D29-FD	0.016	0.092	<d	0.010	0.012
C29	0.258	1.24	0.084	0.050	0.070
D34	1.42	10.6	3.47	0.566	0.691
D34-FD	24.3	203	45.9	1.49	1.49
C34	3.20	26.4	6.58	1.09	1.21
D39	0.067	0.175	0.009	0.049	0.053
D39-FD	0.239	3.31	0.009	0.040	0.038
C39	0.063	0.522	0.009	0.039	0.044
D44	0.108	0.571	0.009	0.347	0.471
D44-FD	0.137	0.540	0.031	0.708	0.950
C44	1.03	8.68	1.54	0.620	0.712
D46	2.52	19.9	3.94	0.711	0.668
D46-FD	0.55	6.54	1.55	0.259	0.282
C46	0.29	1.85	0.16	0.098	0.124

Table 5 (cont'd).					
Field #	Soil concentration (mg/kg)				
	HMX	RDX	TNT	4ADNT	2ADNT
D49	0.092	1.46	0.004	0.015	0.018
D49-FD	0.049	0.166	0.009	<d	0.022
C49	0.062	0.266	0.008	0.026	0.036
D51	34.9	331	81.6	2.23	2.15
D51-FD	5.87	40.9	2.01	1.08	1.10
C51	3.04	21.6	2.68	0.640	0.460
D55	0.517	3.67	1.31	0.108	0.104
D55-FD	0.589	2.68	0.10	0.222	0.274
C55	0.934	8.18	2.52	0.196	0.198
D56	0.043	0.243	<d	0.023	0.027
D56-FD	0.299	2.40	0.350	0.074	0.089
C56	1.05	13.1	0.126	0.066	0.070
D72	0.468	1.40	0.054	0.755	0.813
D72-FD	0.126	0.61	0.036	0.636	0.771
C72	0.968	6.88	1.19	1.07	1.08
D80	0.058	0.187	0.004	0.017	0.031
D80-FD	17.2	179	118	0.081	0.065
C80	0.052	0.266	0.036	0.022	0.036
D82	2.80	24.1	3.29	1.11	1.09
D82-FD	1.22	9.64	0.60	0.67	0.71
C82	0.69	4.50	0.47	0.33	0.33
D83	1.55	7.73	0.783	0.479	0.550
D83-FD	0.76	3.72	0.213	0.248	0.255
C83	0.99	5.48	0.394	0.494	0.564
D87	0.076	0.366	0.003	0.021	0.049
D87-FD	0.071	0.255	0.004	0.071	0.089
C87	0.228	1.504	37.4	0.278	0.260
D90	0.108	0.138	0.004	0.022	0.022
D90-FD	0.035	0.058	<d	0.013	0.016
C90	0.056	0.168	0.190	0.022	0.024

Table 5 (cont'd). Results from the analysis of duplicate discrete and ten-increment composite samples from 20 randomly chosen minigrids near location of low-order mortar detonation. (AcN extraction with RP-HPLC and GC-ECD [shaded] analysis.)

Field #	Soil concentration (mg/kg)				
	HMX	RDX	TNT	4ADNT	2ADNT
D100	<d	0.067	<d	0.013	0.022
D100-FD	1.15	18.8	1.18	0.051	0.063
C100	0.282	2.92	0.982	0.068	0.070
* Minigrid discrete samples.					
† Field duplicate minigrid discrete samples.					
** Ten-increment minigrid composite samples.					

Results for the ten-increment composite samples collected within the same randomly selected minigrids where duplicate discrete samples were collected are also presented in Table 5. These results exhibit the same trends as the field duplicate discrete samples. However, since the composite samples comprise about 5% of the surface area of the 1-m × 1-m minigrid, they should provide a better estimate of the analyte concentration than the discrete samples that comprise only 0.5% of the surface area of the minigrid.

Results for 25-increment composite samples collected within the 10-m × 10-m grid near low-order detonation

Analytical results for the ten random 25-increment composite samples collected within the entire 10-m × 10-m grid are shown in Table 6. The minimum and maximum concentrations for RDX were 4.62 and 294 mg/kg, respectively. This range is only a factor of about 64, whereas the range of concentrations found for the 100 discrete samples from this area differed by nearly five orders of magnitude. However, the relative standard deviation for RDX in these composites was 159% and the median and mean differed by a factor of 2.2, indicating that this group of data for 25-increment composites was not normally distributed. Clearly, very different values can result for random composites, depending on whether or not increments were collected from the apparent hot spots shown in Figure 9. Moreover, because of the presence of a hot spot, energetic residue distribution is clearly not uniform in this grid.

Table 6. Results from the analysis of 25-increment composite samples from grid near a low-order 81-mm mortar detonation at Fort Polk.

Field #	Soil concentration (mg/kg)					Ratio		
	HMX	RDX	TNT	4ADNT	2ADNT	HMX/ RDX	TNT/ RDX	2ADNT/ 4ADNT
G-1-1	4.48	51.0	21.6	0.480	0.608	0.088	0.424	1.27
G-1-2	0.594	4.62	0.752	0.496	0.618	0.129	0.163	1.25
G-1-3	1.02	8.14	1.09	0.484	0.440	0.125	0.133	0.91
G-1-4	32.0	294	106	1.85	1.40	0.109	0.359	0.76
G-1-5	2.14	19.0	4.98	0.422	0.430	0.112	0.262	1.02
G-1-6	8.27	77.9	20.8	0.768	0.791	0.106	0.267	1.03
G-1-7	3.18	25.6	9.70	0.556	0.560	0.124	0.379	1.01
G-1-8	3.30	25.0	5.36	0.660	0.776	0.132	0.214	1.18
G-1-9	2.68	24.0	4.56	0.452	0.516	0.112	0.190	1.14
G-1-10	2.28	16.9	2.76	0.436	0.530	0.135	0.163	1.22
Max	32.0	294	106	1.9	1.4	0.13	0.42	1.3
Min	0.594	4.62	0.752	0.422	0.430	0.088	0.133	0.756
Median	2.93	24.5	5.17	0.490	0.584	0.12	0.24	1.09
Mean	5.99	54.6	17.7	0.660	0.667	0.12	0.26	1.08
Std Dev	*	*	*	0.432	0.284			
% RSD*	*	*	*	65.4	42.7			
<p>* RSDs greater than 100% clearly demonstrate that the data are not normally distributed, therefore, they are not valid statistics.</p> <p>Note: Because the distributions depart greatly from normal as shown by the discrepancy between median and mean, it is not possible to compute useful standard deviations.</p>								

Recent results from sampling at Canadian Force Base–Gagetown (Thiboutot et al. in press) found that multi-increment composite samples collected systematically within a specified area provided reproducible results. We evaluated this approach for this study by creating four mathematical systematic composite samples ($n = 25$) by combining every fourth discrete sample from the 100 discrete minigrids. A comparison of the results for the 100 discrete samples, the ten randomly collected 25-increment composites, and the four systematic mathe-

matal 25-increment composites is shown in Table 7. The range of RDX values is much reduced from a factor of about 10^5 for the discrete samples to factors of 64 and 3, respectively, for the randomly and systematically collected composites. The results for the systematic samples come from only four samples and additional research is needed to verify that this sampling strategy generally provides more reproducible results for this set of environmental conditions. Moreover, it is recognized that the success of a systematic sampling strategy to be reproducible is dependent on the size of the hot spot and the spacing of the sample increments. Likewise, the reproducibility of a random sampling strategy depends on the number of increments. A critical variable for both sampling strategies is the dimensions of the hot spot, which in this case were most likely associated with the partial detonation of an 81-mm mortar.

Table 7. Comparison of concentration estimates for target analytes using various collection strategies in a 10-m × 10-m grid near a low-order detonation.

Collection strategy	Increments per Sample	Number of Replicates	Analytes		
			RDX	HMX	TNT
Discrete samples	1	100			
Max			2390	253	1560
Min			0.037	0.005	0.001
Mean			70.9	7.89	29.7
Median			1.79	0.395	0.044
Composite (random)	25	10			
Max			294	32	106
Min			4.62	0.594	0.752
Mean			54.6	5.99	17.7
Median			24.5	2.93	5.17
Composite (systematic)	25	4			
Max			99.8	10.8	63.2
Min			33.1	3.81	10.3
Mean			70.9*	7.89*	29.7*
% RSD			43.3%	40.0%	77.8%
Median			75.3	8.46	22.7

* Values are the same as the 100 discrete samples.

It is also interesting to compare the median value obtained from the 100 discrete samples with that from the ten 25-increment random and four 25-increment systematic composite samples, because these median values represent

the concentration that half of the sample collected will have lower than (or higher than) values, for this grid. The median of the discrete samples for RDX was 1.79 mg/kg, the median for the ten composite samples was 24.5 mg/kg, and the median for the four systematic samples was 75.3 mg/kg. Thus, rather than diluting out the high concentrations, the multi-increment composite samples are more likely to capture the high concentrations that the discrete samples often miss. A comparison of the discrete and composite (random and systematic) medians for HMX and TNT results in a very similar trend, much higher median concentrations for the composite samples: 0.395 versus 2.93 and 8.5 mg/kg for HMX and 0.044 versus 5.17 and 22.7 mg/kg for TNT, respectively (Table 7).

Table 8. Calculation of total mass of RDX in 10-m × 10-m sampling grid at Fort Polk, Louisiana, and its potential for groundwater contamination.

Grid size = 10 m × 10 m
Grid area = 100 m ²
Sample depth = 0.025 m
Volume of soil sampled = 2.5 m ³ = 2.5E6 cm ³
Soil density = 1.7 g/cm ³
Mass of soil = 1.7 g/cm ³ × 2.5e6 cm ³ = 4.3e6 g = 4.3e3 kg
Weighted average of the average (by total number of increments) RDX concentration for the ten 25-increment random composite samples and the 100 discrete samples 0.059 g/kg
Mass of RDX present in soil to 2.5-cm depth: 0.0059 g/kg × 4.3E3 kg = 250 g
Mass of Composition B collected from grid = 198 g
Composition B is 54% RDX, 6% HMX, 39% TNT, 1% wax.
Mass of RDX in Composition B = 54% of 198 g = 107 g
Total RDX within 100 m ² = 250 g (in soil) + 107 g (in Composition B) = 360 g

A calculation using the weighted average of the average RDX concentration for the ten 25-increment random composite samples and the 100 discrete was made to determine the total mass of RDX present in the topsoil of the 10-m × 10-m grid surrounding the partial detonation event (Table 8). An additional calculation was then made that included both the surface-soil-associated RDX to a depth of 2.5 cm, and the RDX associated with the chunks of Composition B found lying on the surface. The total mass was estimated at 360 g/100 m². By comparison, the 200-m² area sampled around the tank target had a median RDX concen-

tration of 0.948 mg/kg, resulting in a total mass of RDX of 2.04 g/100 m², which is two orders of magnitude lower than the mass in the area surrounding the 81-mm mortar low-order detonation. Even though the target value is smaller than the partial detonation event area, it is three orders of magnitude more than values encountered in previous studies around targets on other artillery impact ranges (Pennington et al. 2001, 2002, 2003). Therefore, although no evidence such as "chunk" energetic residue or large munition fragments was observed, partial detonation events may have occurred near this tank target. In addition, if the partial detonation of an 81-mm mortar was the only source of this energetic residue, then more than one-half of the main charge (560 g RDX) was present on or in the near surface of the 10-m × 10-m grid. Indeed, most of the mass exists as particles that would fit into the size category used for soil (i.e., < 2 mm, Table 8).

Line composite samples surrounding 10-m × 10-m grid

Analytical results for the line composite samples collected at 2-m, 5-m, and 10-m distances off the four edges of the 10-m × 10-m grid are presented in Table 9 and Figure 11. The concentrations of RDX at 2 m off the south and east edges of the grid are 11.3 and 4.56 mg/kg, respectively, which is consistent with the locations where high soil concentrations and particles of solid Composition B were found (Figures 9, 10). The ratios of HMX/RDX and TNT/RDX for the line composite samples are quite similar to those for the discrete minigrid samples, indicating that the contamination is from the same source (i.e., same extent of weathering) Composition B.

For the south, east, and west sides of the grid, concentrations of all analytes decrease as distance from the edge of the grid increases. For the north samples, the highest values for HMX and RDX are found in the 5-m sample. In all cases the samples at a distance of 10 m from the edge are very low compared with most of the concentrations within the grid and the samples collected at 2 and 5 m from the edge. Thus it appears that we have captured the area impacted to the greatest extent from the low-order detonation within the 30-m × 30-m area sampled.

Physical size of hot spot from low-order 81-mm mortar round

In order to develop strategies for hot spot detection, typical physical sizes of these hot spots must be known. It is anticipated that the dimensions of hot spots will depend on both the type of munition and the failure mechanism. Within this 10-m × 10-m area, the major hot spot appears to be centered at minigrid D15. However, since this is based on discrete samples, caution must be stressed, and the subsequent size estimate is tentative. Inspection of Figures 9 and 10 indicates that if we use a concentration of 100 mg/kg as an indicator of the hot spot, the

size of the hot spot around minigrid D15 is about 3 m × 3 m. This is only the first estimate of this type and additional research is planned to provide additional data for dimensions of hot spots from partial detonations of various ordnance items.

Table 9. Concentrations of explosives residues in ten-increment line composite samples collected from the four edges of 10-m × 10-m grid. (Analysis by RP-HPLC and GC-ECD [shaded].)

Edge samples	Soil concentration (mg/kg)						Ratio	
	HMX	RDX	TNT	2,4-DNT	4ADNT	2ADNT	HMX/RDX	TNT/RDX
2 m south	1.39	11.3	2.56	<d	0.444	0.462	0.123	0.227
5 m south	0.316	1.82	0.352	<d	0.144	0.192	0.173	0.193
10 m south	0.076	0.212	0.308	<d	0.130	0.190	0.358	1.453
2 m east	0.614	4.56	1.20	<d	0.149	0.156	0.135	0.262
5 m east	0.180	1.19	0.288	<d	0.070	0.060	0.151	0.242
10 m east	0.098	0.508	0.040	<d	0.030	0.050	0.193	0.079
2 m north	<d	0.130	0.014	0.004	0.020	0.024		0.108
5 m north	0.130	2.25	<d	0.004	0.014	0.016	0.058	
10 m north	0.062	0.198	0.002	<d	0.006	0.008	0.313	0.010
2 m west	0.226	0.930	0.038	0.006	0.072	0.078	0.243	0.041
5 m west	0.134	0.618	0.028	0.004	0.066	0.066	0.217	0.045
10 m west	0.068	0.288	0.020	0.004	0.044	0.046	0.236	0.069
						Mean	0.200	0.248
						Median	0.193	0.108

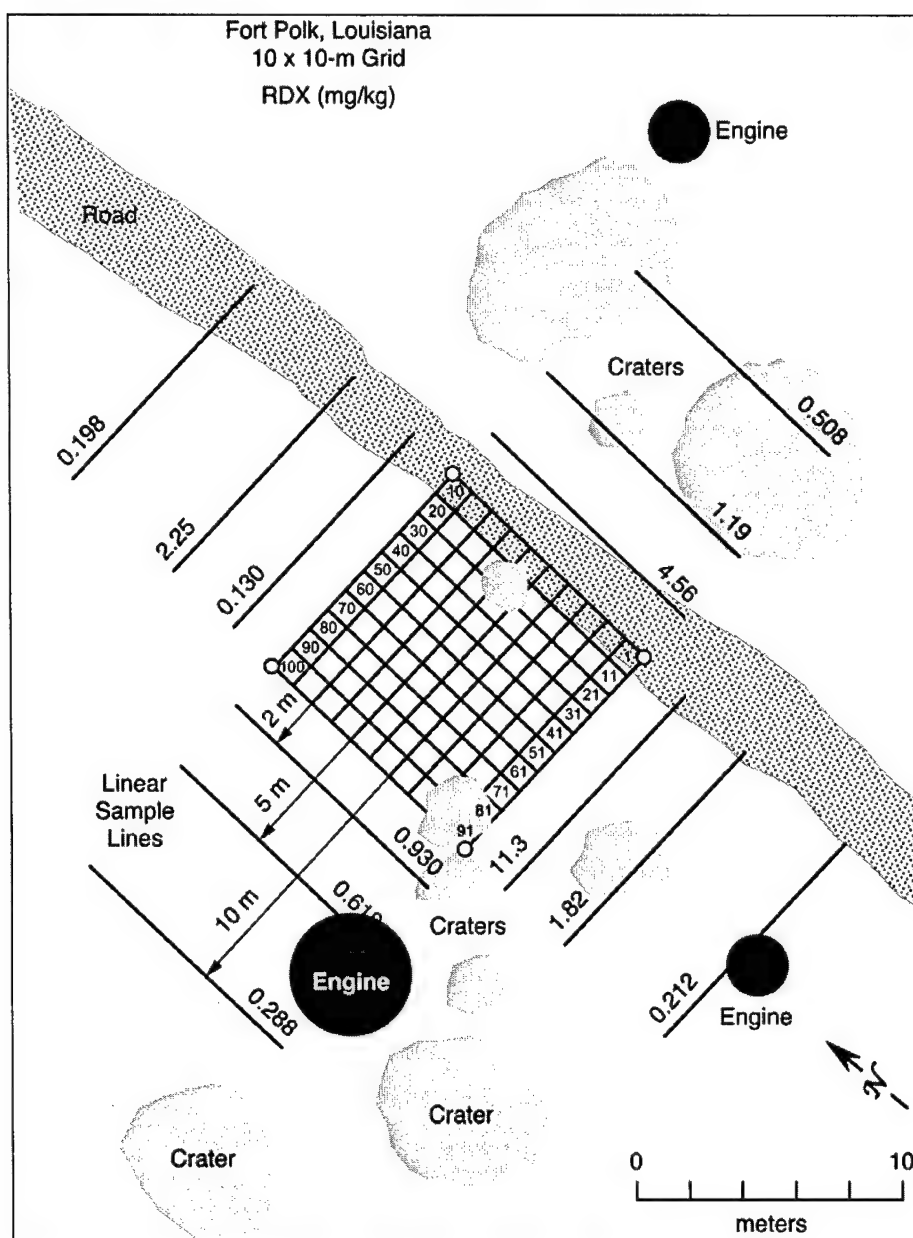


Figure 11. Soil RDX concentrations in linear composite samples taken at various distances from the sampling grid.

Target analyte concentrations near an artillery target

Target analyte concentrations for ten-increment composite surface soil samples that we collected near an artillery target are presented in Table 10 and Figure 12. The concentrations of RDX varied from 0.106 to 15.9 mg/kg, but unlike HMX concentrations near an antitank target (Jenkins et al. 1997, 1998, 2004), there does not appear to be a concentration gradient relative to distance from the target. Therefore, there appears to be a difference in residue pattern around a line-of-sight target as opposed to a target that receives mostly indirect fire. TNT concentrations in these samples varied from 0.076 to 18.8 mg/kg, and the ratios of TNT to RDX were often higher than the 0.7 ratio expected from deposition of fresh Composition B (Table 10). Most 155-mm artillery rounds are filled with TNT rather than Composition B and it appears from the ratio of TNT/RDX that a portion of the explosives residues detected near this target was from TNT-filled rounds. The ratio of HMX to RDX in these samples was also often higher than found in and near the 10-m \times 10-m grid located downhill and to the right of this target. This implies that the Composition B residues near this target are somewhat older (i.e., more weathered) than those near the low-order 81-mm mortar round. This phenomenon occurs because RDX will dissolve faster and migrate away from these residues faster than HMX as a result of its higher aqueous solubility, which exceeds HMX by about a factor of 10.

Table 10. Target analyte concentrations in area around an artillery target in the impact area, Fort Polk.

	Soil concentration (mg/kg)						Ratio		
	HMX	TNB	RDX	TNT	4ADNT	2ADNT	HMX/ RDX	TNT/ RDX	2ADNT/ 4ADNT
NE 0–2 m	15.2	<d	15.9	1.19	0.246	0.308	0.956	0.075	1.25
NW 0–2 m	1.43	<d	1.17	0.144	0.170	0.206	1.22	0.123	1.21
SW 0–2 m	0.420	<d	2.18	0.516	0.280	0.356	0.193	0.237	1.27
SE 0–2 m	0.360	0.082	0.500	18.8	0.912	1.17	0.720	37.7	1.28
ENE 2–5 m	0.876	<d	0.448	0.436	0.172	0.230	1.96	0.973	1.34
NNE 2–5 m	0.236	<d	0.724	0.076	0.074	0.096	0.326	0.105	1.30
NNW 2–5 m	0.216	<d	1.75	14.5	0.266	0.248	0.123	8.24	0.93
WNW 2–5 m	0.120	<d	0.422	0.234	0.176	0.274	0.284	0.555	1.56
WSW 2–5 m	1.92	<d	13.3	4.42	0.526	0.732	0.144	0.333	1.39
SSW 2–5 m	0.225	<d	1.23	2.22	0.611	0.875	0.183	1.81	1.43
SSE 2–5 m	0.134	<d	0.294	9.46	1.14	1.42	0.456	32.2	1.25
ESE 2–5 m	0.064	<d	0.106	0.782	0.296	0.400	0.604	7.38	1.35
						Mean	0.597	7.47	1.30
						Median	0.391	0.764	1.29

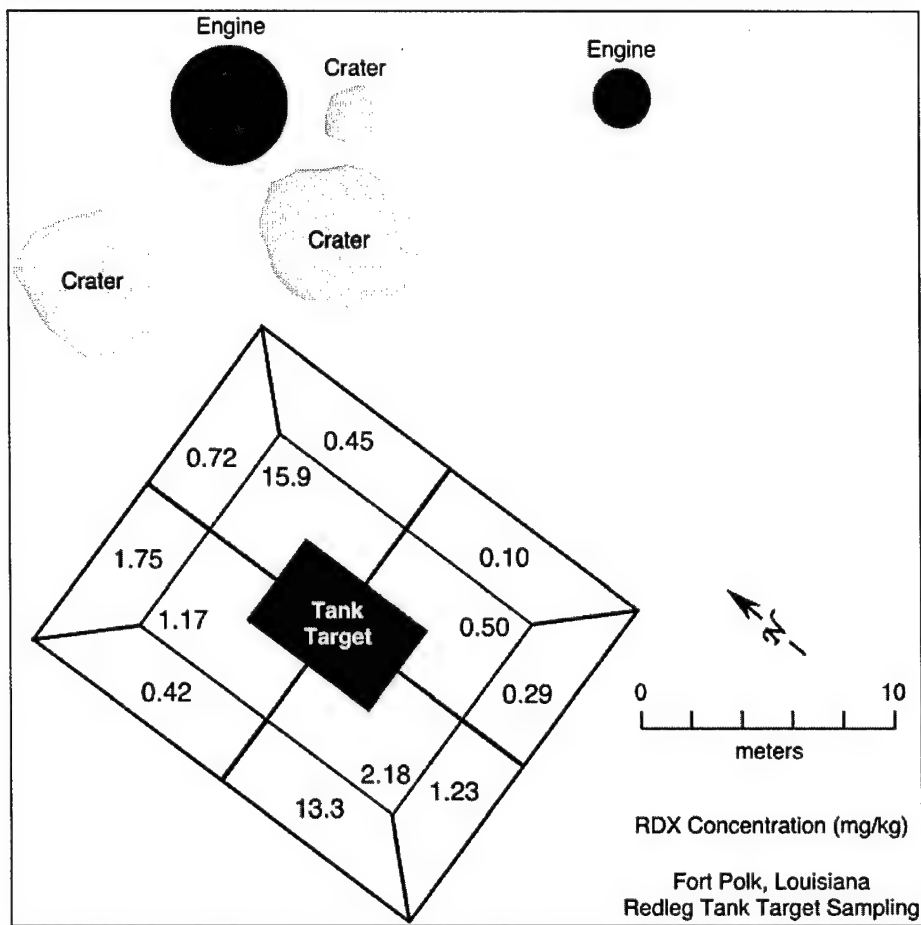


Figure 12. Soil concentration of RDX taken in various sampling areas around a tank target.

5 SUMMARY AND CONCLUSIONS

Surface soil samples were collected and analyzed for explosives residues from the artillery/mortar impact area located at Fort Polk, Louisiana. Two distinct areas were selected for sampling. The first was around a low-order detonation event and the second around a tank target.

The first sampling area was selected as a result of the observation of numerous pieces of Composition B residue lying on the soil surface, thereby giving us the opportunity to sample a potential "hot spot." A 10-m \times 10-m sampling grid that encompassed the residue chunks of Composition B was laid out. This grid was further subdivided into 100 1-m \times 1-m minigrids. Observed pieces of Composition B were collected and weighed and their position within the grid system was cataloged. One discrete soil sample was collected from each of the minigrids. The most predominant analyte, RDX, had concentrations ranging over approximately five orders of magnitude for the 100 discrete samples. TNT concentrations ranged from less than detect (0.002 mg/kg) to 1560 mg/kg or six orders of magnitude. Certainly no single discrete sample could accurately represent the entire grid area.

Field duplicate discrete samples were collected from 20 randomly chosen minigrids. The differences between these field duplicates varied up to three orders of magnitude, indicating that single samples cannot represent areas as small as one minigrid (square meter) for energetic materials that exist as crystalline particulates. Ten-increment composites were also collected within these twenty minigrids. There was no apparent agreement between the initial discrete and the composite samples, indicating again that the discrete sampling method does not represent the area of concern for RDX, TNT, and HMX. Ten composite samples of 25 randomly chosen increments each were taken over the entire 10-m \times 10-m grid. The median RDX concentration for the ten composite samples was fourteen times higher than the median of the 100 discrete samples because the "hot spots" were more frequently sampled with the former sampling strategy than were the latter. Also, RDX concentrations for these composite samples varied as much as 60 times as a result of the number of times the "hot spot" was sampled. Therefore, under these conditions, much uncertainty exists among composite samples composed of 25 randomly collected increments, even though a large improvement over discrete sampling was achieved.

We decided to mathematically generate systematic random samples by "compositing," from the 100 discrete samples, every fourth minigrid. We did this four times, thereby using all of the minigrid samples. The median value for RDX was

three times greater than the ten random composites median and 42 times the median for the discrete samples. The RDX concentration range for the results of these four systematic mathematical composites was 33 to 100 mg/kg, whereas the range was 5 to 294 mg/kg, and 0.04 to 2,390 mg/kg for the ten 25-increment randomly collected composites and the 100 discrete samples, respectively. It appears the systematic approach is more likely to consistently sample "hot spots" of the size encountered in this study.

Because most of the discrete samples had concentrations well below the mean, we divided the results into those less or greater than 100 mg/kg in order to delineate the area of highest concentration. The majority of these later samples were co-located within an area of approximately 3 m \times 3 m. The remaining high values were from two diagonally joined minigrids.

Composite soil samples were also collected along linear transects at 2, 5, and 10 m from the grid on all four sides. RDX concentrations decreased with distance from the grid to less than 0.3 mg/kg at the 10-m distances. It appears that this detonation event extended its influence across an area of approximately 30 m \times 30 m if one chose a boundary concentration of 0.3 mg/kg.

The second sampling area, a tank target, was selected because of the fact that it would be a point of interest for incoming fire. A sampling scheme different from that used to delineate the area of influence of the low-order detonation event was used at the target. A two-meter-from-the-target grid was set around the tank and then quartered. An additional grid was set at 5 m from the tank and this 2- to 5-m area was divided into eight parts. Ten-increment composite soil samples were collected from inside each of the twelve areas. RDX concentrations varied from 0.1 to 16 mg/kg. Within the described target sampling area there was no apparent pattern to the distribution of RDX, i.e., there was no concentration gradient moving out from the target, such as those found around targets at anti-tank ranges. One might anticipate this, as direct fire from firing point to target is used at antitank ranges. This means that the munition comes from one direction and usually impacts the target. Any casing rupture or partial detonation would be at the target and the majority of HE would be deposited there and spattering would decrease with distance from the target. At artillery and mortar impact areas, the HE round can arrive from the air from numerous directions, frequently from all around the target. Partial detonation events could occur anywhere around the target and possibly at considerable distance from the target, depending on the accuracy of the gunner.

This study reinforces earlier work, the results of which indicated that low-order (partial) detonation events produce the most HE residues within impact areas. In most cases pure HE material is found and residue concentrations in the

soil are higher than the surrounding area. This work was a first attempt to delineate the area of influence of one of these events. Similar studies need to be made to provide more estimates of the areas influenced by the low-order detonations of other munitions. Additionally, systematic random composites seem to produce a more reproducible and regulatory appropriate sample than that generated by a random composite, and certainly either provides results that are an improvement over discrete sampling methods. Additional field studies are needed to compare systematic random composites with random composites under a variety of conditions and events. Also, sampling studies need to be conducted over larger areas than the 10-m \times 10-m area studied here.

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